

Pechkovskii, V. V.

Thermochical decomposition of sulfates in the presence  
of copper. II. Experimental part. V. V. Pechkovskii.

Nauk. Trudy, Nauchnitsk. Politekhn. Inst. im. S.

Oroshenikisza 1955, No. 30, 292-3; cf. U.S. Pat. 2,904,111.

The rate of reaction and the products formed by the thermal  
decomposition of  $\text{CuSO}_4$ ,  $\text{ZnSO}_4$ ,  $\text{MnSO}_4$ , and  $\text{MgSO}_4$  in the  
presence of Cu were investigated. Expts. showed that Cu  
was oxidized with gaseous O formed by the dissociation of  
sulfates.

T. Darbala

PECHKOVSKIY, V V

Reaction of amides with metal oxides. M. R. Pozin.  
A. M. Ginstrom and V. V. Pechkovskiy. J. Appl. Chem.  
U.S.S.R. 28, 611-15 (1955) (Engl. translation).—See C.A.  
50, 101.

B. M. R.

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(2)

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AID P - 2276

Subject : USSR/Chemistry

Card 1/1 Pub. 152 - 2/21

Author : Pechkovskiy, V. V.

Title : Study of the thermal decomposition of manganese sulfate

Periodical: Zhur. prikl. khim., 28, no.3, 237-244, 1955

Abstract : The mechanism of the decomposition of both pure manganese sulfate and a mixture of manganese sulfate with carbon has been studied. Almost pure Mn<sub>3</sub>O<sub>4</sub> was obtained by conducting the decomposition of MnSO<sub>4</sub> at 800°C. Six tables, 4 diagrams, 9 references (7 Russian: 1937-52).

Institution: Novocherkassk Polytechnic Institute (im. Sergo Ordzhonikidze)

Submitted : 0 19, 1953

AID P - 3722

Subject : USSR/Chemistry  
Card 1/1 Pub. 152 - 2/16  
Author : Pechkovskiy, V. V.  
Title : Study of conditions of sulfating roasting of copper sulfide in order to obtain copper sulfate  
Periodical : Zhur. prikl. khim. 28, 8, 802-806, 1955  
Abstract : Addition of cupric oxide to copper sulfide accelerates the formation of sulfate from sulfide. At 500°C, 26% sulfide was converted to sulfate without addition of cupric acid (in 60 min.). In the presence of 25% CuO, 39% CuSO<sub>4</sub>, and in the presence of 50%, 47.5% CuSO<sub>4</sub> were obtained resp. under identical reaction conditions. Three tables, 3 diagrams, 3 references, all Russian (1938-1949).  
Institution : Novocherkassk Polytechnic Institute im. Sergo Ordzhonikidze  
Submitted : Dec. 18, 1954

PECHKOVSKIV VV

Recovery of potassium and magnesium sulfate by thermal decomposition of langbeinite. V. V. Pechkovskii and N. A. Iakhskova (Tsvetnochim. Inst., Novosibirsk). Zhur. Tsvet. Khim., 11, 1166-69 (1966).—Mixts. of langbeinite ( $\text{NaCl}$  free) with C were heated in a current of air. The percentage of total S (based on  $\text{K}_2\text{SO}_4$ ) passing into the gaseous phase increased linearly with the temp. and increased with the C content in the mixt. The percentage of  $\text{SO}_2$ , %, in the gaseous phase increased linearly with the temp. up to 66% at 900° and then decreased to 16.2% at 1100°; simultaneously, the percentage of elemental S, %, increased continuously to 33.7% at 950°.  $k'$  and  $k''$  are functions of the temp. and the C content as follows: at 500, 800, 900, 950, and 1100° in mixts. contg. 15% C,  $k' = 5, 42.0, 62.8, 18.0$ , and  $66.3$  and  $k'' = 1.1, 2.8, 5.2, 12.5$ , and 32.7%; with 20% C,  $k' = 27.0, 41.9, 59.2, 62.1$ , and 78.8%; with 25% C,  $k' = 1.5, 3.0, 6.0, 37.8$ , and 78.8%; with 25% C,  $k'' = 32.0, 58.7, 61.2, 78.1, \dots$ , and  $k'' = 1.8, 3.4, 24.6, 4.0, \dots$ ;  $k''$  increased rapidly during the 1st 10 min. and tended to become const. I. Beinowitz

PECHKOVSKY, V.V.

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*V. Oxidation of sulphides and sulphide-ores. M. E. Nuzin, A. M. Gavrilov and V. V. Pechkovskiy (Zh. prikl. Khim., 1955, 28, 1249-1254).—The temperature at which intensive evolution of  $SO_2$  begins when air is passed over Cu, ZnS and CuS rises by this order, and is very close to that for sulphate-sulphide mixtures in a stream of  $N_2$ . This suggests the after-initial oxidation of surface sulphide. The velocity of the reaction between sulphide and sulphate determines the rate of  $SO_2$  evolution, and is itself controlled by the velocity of diffusion of  $O_2$  and  $SO_4^{2-}$  through the surface layer. At a given temp., production of  $SO_2$  from mixtures of sulphides exceeds that found with each sulphide taken separately. This effect is related to differences in the relative rates of oxidation of the sulphides and of decompr. of the resulting sulphates, as a result of which the sulphido-one metal is able to react with the sulphate of another. R. Ruscon.*

*PECHKOVSKII, V. V.*

PECHKOVSKII, V. V., dots., kand. khim. nauk.

Interaction between molybdenum trioxide and some metal oxides.  
Trudy NPI 27:167-171 '56. (MIRA 10:12)

1. Kafedra tekhnologii neorganicheskikh veshchestv Novocherkasskogo  
politekhnicheskogo instituta.  
(Metallic oxides)

*PECHKOVSKY, V. V.*

Reaction of carbon dioxide with metal oxides in the presence of sulfur dioxide. I. (A. M. Gorkov State University, Novosibirsk). ZH. FIZ. Khim. 30, 2352 (1957) p. 69, 1137 (B. 50, 10330). - Metal oxides with and without C were heated in a current (1 l/min.) of 10% SO<sub>2</sub> gas for 15 min. The exp'tl. results were correlated with thermodynamic data from partial pressures of the gaseous products. Without C the rate of sulfate formation X with Li<sub>2</sub>O, CaO, and Mn<sub>2</sub>O<sub>3</sub> increased with the temp. and passed through a max. at 400, 500, and 600°, resp. In the same temp. range (500-700°) with MgC X increased continuously. The results supported the reaction mechanism: MO + SO<sub>2</sub> → MSO<sub>4</sub>; MSO<sub>4</sub> + 1/2 O<sub>2</sub> → MSO<sub>4</sub>, and MO + 2M + 1/2 O<sub>2</sub>. In mixt. with C the value of X increased with the temp., and mol. ratio C/MO = M. For the same values of M the values of X, i.e. different oxides, increased in the following order: Al<sub>2</sub>O<sub>3</sub>, MgO, Fe<sub>2</sub>O<sub>3</sub>, ZnO, Mn<sub>2</sub>O<sub>3</sub>, Co<sub>3</sub>O<sub>4</sub>, and Cu<sub>2</sub>O. Apparently the reaction of SO<sub>2</sub> with MO in the presence of C is controlled either by the rate of reduction of MO or by the rate of MS formation. The absorption of S by C increased with the temp. and was held as tentatively that it could not be exst. with CS<sub>2</sub>.

"APPROVED FOR RELEASE: 06/15/2000

CIA-RDP86-00513R001239810018-8

35-2451-2, Vol. 1, Part IV, Item 2.

Apparatus for low temperature study of the properties of precision  
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(MIRA 1945)

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CIA-RDP86-00513R001239810018-8"

ACCESSION NR: AP4034578

S/0076/64/038/004/0916/0920

AUTHOR: Amirova, S. A. (Perm'); Pechkovskiy, V. V. (Perm'); Prokhorova, V. G. (Perm'); Ostrovskaya, T. V. (Perm'); Lezhneva, A. A. (Perm')

TITLE: Oxidation of iron-vanadium spinel by oxygen.

SOURCE: Zhurnal fizicheskoy khimii, v. 38, no. 4, 1964, 916-920

TOPIC TAGS: iron vanadium spinel, oxidation, thermogram, iron orthovanadate containing system, vanadium pentoxide containing system, iron orthovanadate, solid subtraction solution, vanadium hematite solution, fusion temperature, solubility, alkali additive, oxidation acceleration

ABSTRACT: This investigation of the oxidation of iron-vanadium spinel by oxygen included a study of the composition and properties of the phases formed, and the effect of small amounts of alkali additives on the oxidation process. Thermograms for the iron-vanadium spinel system, for iron orthovanadate and for the iron orthovanadate-vanadium pentoxide system were constructed. In the oxidation of the spinel the formation of a solid subtraction solution (exotherm at 236-336°C, spinel crystal structure is retained but the cell parameters decreased) proceeds

Card 1/2

ACCESSION NR: AP4034578

decomposition of the spinel. Three phases are formed by the oxidation of the spinel (exotherm at 462-573°C): vanadium pentoxide, iron orthovanadate and the phase  $R_2O_3$  which represents a solid solution of vanadium in hematite. The endo-therms 619-641 and 790-850°C correspond to the fusion of the eutectic of vanadium pentoxide and iron orthovanadate and the pure  $FeVO_4$ . The solubility of iron orthovanadate in 10%  $H_2SO_4$  was determined. The addition of 0.5% KCl to the spinel greatly accelerates its oxidation but does not affect the oxidation products.  
Orig. art. has: 2 tables and 4 figures.

ASSOCIATION: Permskiy politekhnicheskiy institut (Permsk Polytechnical Institute)

SUBMITTED: 28Apr63

ENCL: 00

SUB CODE: MM, GC

NO REF SOV: 003

OTHER: 001

Card 2/2

PECHKOVSKIY, V. V.

V Thermocchemical decomposition of aluminum sulfate  
V. V. Pechkovskiy and A. N. Kulin (A. M. Gorki State  
Univ., Majotov). Zhur. Priklad. Khim. 70, 1500-10  
(1987). — The percentage decompr. ( $\alpha$ ) of  $\text{Al}_2(\text{SO}_4)_3$  was  
determ. by the loss in wt., and the total S in the gases was  
detd. as  $\text{SO}_2$  by preliminary reduction of all the gases formed  
with CO at 150-750°.  $\alpha$  increased with the temp. and duration  
(15-90 min.). The values of  $\alpha$  (30 min.) in dry air at  
750, 800, 840, and 900° were 8.8, 12.0, 33.1, and 87.4%,  
resp. The addn. of 5%  $\text{Fe}_2\text{O}_3$ ,  $\text{Cr}_2\text{O}_3$ , or  $\text{TiO}_2$  increased  
these values to 8.7, 9.1, and 7.1% at 750°; to 18.6, 20.8,  
and 11.1% at 800°; and to 7.1, 33.0, and 30.0% at 850°.  
Increasing the proportion of added oxides from 5 to 10% did  
not affect  $\alpha$ . At 800° in a current of CO in 15 min.,  $\text{Al}_2(\text{SO}_4)_3$  was decomprd. to the extent of 78.4% with formation  
of  $\text{Al}_2\text{O}_3$  and a gas contg. 70.3%  $\text{SO}_2$  and 8.1% of  $\text{CO}_2 + \text{CS}_2 + \text{S}_2$ .

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+Pechkovskiy V.V.

Reaction of sulfur dioxide with metal oxides in an oxidizing atmosphere. V. V. Pechkovskii (In. M. Gorki State Univ., Leningrad), Zhur. Priklad. Khim.-40, 1679-82(1967); J. C. A. 51, 10067A. - Powd. metal oxides were heated in a current of dry 2.5-20% SO<sub>2</sub>. The degree of conversion  $\eta$  to sulfate of Fe<sub>2</sub>O<sub>3</sub> and CuO passed through a max. at 800° and of Cr<sub>2</sub>O<sub>3</sub> at 800°, whereas  $\eta$  of Mn<sub>2</sub>O<sub>3</sub>, MgO, and CaO increased continuously with the temp. (1000°).  $\eta$  increased with the concn. of SO<sub>2</sub>. The presence of (9%) Cr<sub>2</sub>O<sub>3</sub>, Fe<sub>2</sub>O<sub>3</sub>, CuO, Cr<sub>2</sub>O<sub>3</sub>, or Mn<sub>2</sub>O<sub>3</sub> catalyzed the conversion of MgO. In these mixts. sulfates of Fe and Cu were not formed, whereas sulfates of Co and Mn were present with that of MgO. A relation between  $\eta$  and the degree of "likeness" of a given oxide to SO<sub>2</sub>, i.e. the thermal stability of the sulfate, was noted. Thus in the order of increasing  $\eta$  the following oxides are arranged: MnO < MgO < CaO and Fe<sub>2</sub>O<sub>3</sub> < CuO < Cr<sub>2</sub>O<sub>3</sub> < Mn<sub>2</sub>O<sub>3</sub>. The catalytic effect on  $\eta$  of MgO is in the decreasing order Cr<sub>2</sub>O<sub>3</sub> > Fe<sub>2</sub>O<sub>3</sub> > CuO. 1. Berezovitz

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Pechkovskiy, V. V.

USSR/Physical Chemistry - Thermodynamics, Thermochemistry, Equilibria,  
Physical-Chemical Analysis, Phase Transitions.

B-8

Abs Jour: Referat. Zhurnal Khimiya, No 2, 1958, 3753.

Author : V.V. Pechkovskiy.

Inst :  
Title : Thermochemical Dissociation of Zinc Sulfate.

Orig Pub: Zn. neorgan. khimii, 1957, 2, No 7, 1467-1470.

Abstract: The speed of the thermal dissociation of  $ZnSO_4$  and the composition of solid and gaseous calcination products depending on the temperature, experiment duration and  $O_2$  concentration in gas flowing over the heated sulfate were investigated. The  $ZnSO_4$  dissociation in a flow of air proceeds at a noticeable speed even at  $750^\circ$  with formation of  $ZnO$ ,  $SO_2$ ,  $SO_3$  and  $O_2$ . With an increase of temperature and experiment duration and a decrease of  $O_2$  concentration, the  $SO_3$  content in the calcination gas is dropping (especially at  $850$  to  $900^\circ$ ), but it always re-

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Card : 2/2

PECHKOVSKIY, V.V.

Sulfating manganese compounds with sulfur dioxide. Zhur. prikl.  
khim. 29 no.7:977-980 Jl '57. (MIRA 10:10)

*Perer*  
1. Moskovskiy gosudarstvennyy universitet im. A.M. Gor'kogo.  
(Manganese compounds) (Sulfur dioxide)

*Pechkovskiy et al.*  
AMIROVA, S.A.; PECHKOVSKIY, V.V.; PARKACHEVA, V.V.

Studying the roasting of carbonaceous pyrite. Zhur.prikl.khim.  
30 no.12:1735-1740 D '57. (MIRA 11:1)

1.Permskiy gosudarstvennyy universitet imeni A.M. Gor'kogo.  
(Pyrites)

SOV/156-58-3-42 17

## AUTHORS:

Fechkovskiy, V. V., Amirova, S. A., Parkacheva, T. V.

## TITLE:

The Investigation of the Sulfatization by the Combustion of  
the Cobalt Concentrates in a Pseudo-Boiling Charge  
(Issledovaniye sulfatiziruyushchego obzhiga kobaltovogo  
kontsentrata v psevdozhizhennom sloye)

## PERIODICAL:

Nauchnyye doklady vysshey shkoly, Khimiya i khimicheskaya  
tekhnologiya, 1958, Nr 3, pp. 592-594 (USSR)

## ABSTRACT:

In the present paper the sulfatization of cobalt concentrate  
in combustion in the laboratory was investigated. The degree  
of sulfatization of the cobalt; the sulfur combustion; the  
smoke separation and its dependence on the temperature; and  
the composition of the concentrate and the granular size of  
the particles in the concentrate were determined. It was found  
that the smoke separation in the combustion of cobalt con-  
centrates in the boiling charge of the laboratory furnace  
depends on the air velocity, the physical and chemical  
properties of the material, the temperature, and the granular  
state of the material. At 450°C the smoke separation amounts  
to 2 - 3 %, at 600-700°C to 16 - 20 %. Also, the separati

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SOV, 156-58-3-49/52

The Investigation of the Sulfatization by the Combustion of the Cobalt Concentrates in a Pseudo-Boiling Charge

of sulfur in the combustion of cobalt concentrate is dependent on the temperature and the duration of the combustion, the thickness of the charge of the material to be burned, the chemical composition and the granulation. The degree of sulfatization of the cobalt also depends on these same factors. Above 550° the sulfatization of cobalt decreases. The formation of cobalt sulfate is increased when a great part of the sulfidic sulfur of the material has already been burned. According to the material collected experimentally the sulfatization process takes place in two stages:

- 1) The oxidation of sulfides.
- 2) The formation of sulfates.

There are 1 figure, 1 table, and 6 references, 5 of which are Soviet.

Card 2/3

The Investigation of the Sulfatization by the Combustion of the Cobalt Concentrate in a Pseudo-Ballistic Charge

SGV/156-58-3-47/12

ASSOCIATION: **Kafedra** tehnologii neorganicheskikh veshchestv  
Leningradskogo gosudarstvennogo universiteta im. A. M. Gor'kiy  
(Chair for Technology of Inorganic Substances, Leningrad State University imeni A. M. Gor'kiy)

SUBMITTED: January 24, 1958

Card 5/5

**AUTHORS:**

Ketov, A. N., Pechkovskiy, V. V.

SOV/156-58-4-14/49

**TITLE:**On the Mechanism of the Thermal Decomposition of the Sulfites  
of Calcium, Strontium and Barium (O mekhanizme termicheskogo  
razlozheniya sul'fitov kal'tsiya, strantsiya i bariya)**PERIODICAL:**Nauchnyye doklady vysshey shkoly. Khimiya i khimicheskaya  
tekhnologiya, 1958, Nr 4, pp 667-671 (USSR)**ABSTRACT:**

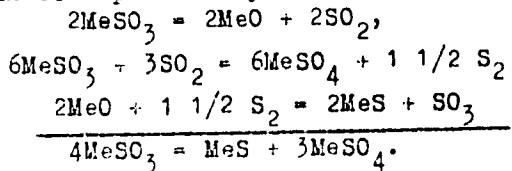
In the present paper investigations were carried out concerning the thermal decomposition of the sulfites of calcium, strontium and barium. This was done to explain the mechanism of this thermal decomposition. The composition of the solid and gaseous products of the above-mentioned sulfites were determined, as well as the decomposition in dependence on temperature, time of stay and the type of atmosphere. The investigations were carried out with specially synthesized and purified anhydrous sulfites. SO<sub>2</sub> and elementary sulfur were determined in the gaseous phase. In the solid phase sulfur was determined in the form of sulfite, sulfide, thiosulfate and sulfate. Within the temperature range of from 600° to 800°C no melting of the sulfites and their decomposition products occurs. The process

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SOV/156-58-4-14/49

On the Mechanism of the Thermal Decomposition of the Sulfites of Calcium,  
Strontium and Barium

of thermal decomposition of sulfites of calcium, strontium and  
barium can be expressed by the following equation:



The formation of the sulfites and sulfates takes place under  
direct action of the gaseous phase. The density of sulfates  
formed in the decomposition exerts a decisive influence upon  
the rate of thermal decomposition of the sulfites of calcium,  
strontium and barium. The dependence of thermal decomposition  
upon the time of stay in nitrogen current at 700° after 5, 10  
and 15 minutes was investigated, and it was found that the  
strontium sulfite is decomposed most easily.  
There are 3 tables and 5 references, 4 of which are Soviet.

ASSOCIATION: Kafedra tekhnologii neorganicheskikh veshchestv Permskogo  
Card 2/3 gosudarstvennogo universiteta im. A. M. Gor'kogo (Chair of the

SOV/156-58-4-14/49

On the Mechanism of the Thermal Decomposition of the Sulfites of Calcium,  
Strontium and Barium

Technology of Inorganic Substances at the Perm' State University  
imeni A. M. Gor'kiy)

SUBMITTED: May 24, 1958

Card 3/3

AUTHOR: Pechkovskiy, V.V. SOV/ 78-3-7-3/44

TITLE: On the Thermodynamic Decomposition of Zinc Sulfate in the Presence of Oxides of Iron, Chromium, Copper, and Aluminum  
(O termicheskom razlozhenii sul'fa a tsinka v prisutstvii okisley zheleznykh, khroma, merti i alumininiy)

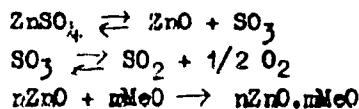
PERIODICALS: Zhurnal neorganicheskoy khimii, 1958, Vol 3, Nr 7, pp 1483-1486  
(USSR)

ABSTRACT: The velocity of the thermal decomposition of zinc sulfate in the presence of some metal oxides such as iron-, chromium-, copper-, and aluminum oxide was investigated. It follows from the experiments carried out that the thermal decomposition of zinc sulfate is only little accelerated by the addition of  $Al_2O_3$ , but that, by the addition of  $Cr_2O_3$ ,  $CuO$ ,  $Fe_2O_3$ , i.e. in the case of such additions in which metals show a change of valence, the thermal decomposition of zinc sulfate is accelerated. The thermal decomposition of zinc sulfate in the presence of metal oxides develops according to the following equations:

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On the Thermodynamic Decomposition of Zinc Sulfate in  
the Presence of Oxides of Iron, Chromium, Copper, and  
Aluminum

SOV/ 78-3-7-3/44



The acceleration of the thermal decomposition of zinc sulfate by the addition of  $\text{Cr}_2\text{O}_3$ ,  $\text{Fe}_2\text{O}_3$ , and  $\text{CuO}$  is based upon the catalytic action brought to bear upon zinc sulfate by these oxides. The best catalytic effect is produced by  $\text{Cr}_2\text{O}_3$ , while that of  $\text{CuO}$  and  $\text{Fe}_2\text{O}_3$  is not so strong. There are 3 figures, 3 tables, and 5 references, which are Soviet.

SUBMITTED: June 26, 1957

1. Zinc sulfate--Decomposition
2. Zinc sulfate--Thermal effects
3. Aluminum oxide--Chemical reactions
4. Copper oxides--Chemical reactions
5. Iron oxides--Chemical reactions

Cart 2/2

PACHKOVSKIY, V.V.

Decomposition of zinc and cobalt sulfates in a stream of sulfur  
dioxide - air mixture. Zhur. prikl. khim. 31 no.8:1139-1143 Ag '58.  
(MIRA 11:10)

1. Permskiy gosudarstvennyy universitet imeni A.M. Gor'kogo.  
(Zinc sulfate) (Cobalt sulfate)

FECHKOVSKIY, V.V.; AMIROVA, S.A.; PARKACHEVA, V.V.

Obtaining valuable components from flotation pyrite by sulfating  
roasting. Z'ur. prikl. khim. 31 no.10:1466-1471 O '58.  
(MIRA 12:1)

1.Pernskiy gosudarstvennyy universitet imeni A.M. Gor'kogo.  
(Pyrites) (Metallurgy)

KETOV, A.N.; PECHKOVSKIY, V.V.

Reaction of sulfur dioxide with calcium oxide and calcium carbonate. Zhur.prikl.khim. 31 no.12:1783-1787 D '58. (MIRA 12:2)

1. Permskiy gosudarstvennyy universitet imeni A.M. Gor'kogo.  
(Sulfur dioxide) (Calcium oxides) (Calcium carbonates)

SOV/156-59-2-46/48

15(2)

AUTHORS:

Amirova, S. A., Pechkovskiy, V. V., Prokhorova, T. G.,  
M. I. Polotnyanshchikova

TITLE:

The Examination of the Oxidizing and Chlorinating Burning of  
Vanadium-Slag (Izuchenie okislitel'nogo i khloriruyushchego  
obzhiga vanadiyevogo shlaka)

PERIODICAL:

Nauchnyye doklady vyschey shkoly. Khimiya i khimicheskaya  
tekhnologiya, 1959, Nr 2, pp 398-401 (USSR)

ABSTRACT:

The production of vanadium from converter-slag by oxidizing  
burning with sylvinitite, potassium-, or sodiumchlorite or by  
treatment with dry chlorine gas is investigated. Finely crushed  
slag was mixed with various admixtures and burned in a  
laboratory furnace under a stream of air or chlorine. The  
portion of soluble vanadinites which had formed after the  
burning, was analytically determined. When treated with chlorine,  
the waste gas was condensed, and the content of V, Fe, and  
Ti was determined in the condensate. The results are shown in  
(Tables 1-3). The best yield of vanadium is obtained at  
temperatures of from 800 to 850 degrees. Higher temperatures  
caused overbaking and thereby reduced the yield. The use of  
sylvinitite, potassium-, or sodiumchloride made no differences

Card 1/2

The Examination of the Oxidizing and Chlorinating  
Burning of Vanadium-Slag

S07/156-59-2-46/48

in the yield, but the reaction develops faster with the two potassium salts. Fine-grain crushing of the slags results in higher yields. The yield of vanadium and iron by chlorinating the slags is shown in table 4. The ironchlorides condensate much easier than the vanadium chlorides. The mixture of iron- and vanadium-chlorides could therefore be separated by distillation. The addition of carbon increases the yield of chlorides, but leads to the forming of volatile titanium-chlorides, which pass into the condensate. There are 4 tables and 9 Soviet references.

PRESENTED BY: Kafedra tekhnologii neorganicheskikh veshchestv Parmskogo gosudarstvennogo universiteta im. A. M. Gor'kogo (Chair for Technology of Inorganic Materials Perm' State University imeni A. M. Gor'kiy)

SUBMITTED: December 29, 1958

Card 2/2

PECHKOVSKIY, V.V.

Sulfidizing process during the reduction of calcium sulfate.  
Izv.vys.ucheb.zav.; tevet.met. 2 no.4:54-58 '59.  
(MIRA 13:1)

1. Permskiy gosudarstvennyy universitet. Kafedra tekhnologii  
neorga nicheskikh veshchestv.  
(Nickel--Metallurgy) (Calcium sulfate)

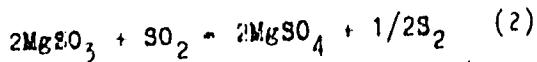
SOV/78-4-2-5/40

. 5(4)  
AUTHORS: Ketov, A. N., Pechkovskiy, V. V.  
TITLE: Investigation of the Thermal Decomposition of Magnesium Sulfite  
(Izucheniiye termicheskogo razlozheniya sul'fita magniya)  
PERIODICAL: Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 2,  
pp 272-276 (USSR)  
ABSTRACT: The velocity of the thermal decomposition of magnesium sulfite  
depending on the temperature, the type of the gaseous re-  
ducing agent, and on various additions as  $\text{SiO}_2$ ,  $\text{Fe}_2\text{O}_3$ ,  $\text{Cr}_2\text{O}_3$ ,  
and ferriferous bauxite, were investigated. Sulfur dioxide,  
sulfur, magnesium thiosulfate, and magnesium sulfate are  
formed in nitrogen atmosphere in the temperature range of  
300-500°. At temperatures  $> 500^\circ\text{C}$  only sulfur dioxide,  
elementary sulfur and magnesium sulfate are formed. The forma-  
tion of sulfur dioxide during the thermal dissociation of  
magnesium sulfite proceeds according to the following reaction:  
$$\text{MgSO}_3 \rightleftharpoons \text{MgO} + \text{SO}_2 \quad (1)$$
  
The formation of elementary sulfur and of magnesium sulfate  
proceeds according to the following reaction:

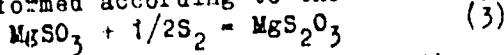
Card 1/4

SOV/78-4-2-5/40

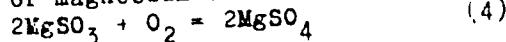
Investigation of the Thermal Decomposition of Magnesium Sulfite



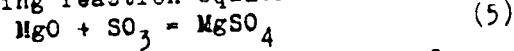
At temperatures < 500° and in a nitrogen atmosphere magnesium sulfite reacts, during magnesium thiosulfate formation, with the sulfur being formed according to the following equation:



On a rise of the oxygen concentration in the gaseous phase the formation of magnesium sulfate increases:



On an addition of iron and chromium oxide at 500° a quicker decomposition of magnesium sulfite takes place. The SO<sub>2</sub> formation decreases because SO<sub>3</sub> is formed under the catalytic influence of the oxides. This SO<sub>3</sub> reacts with magnesium oxide, in connection with the formation of magnesium sulfate, according to the following reaction equation:



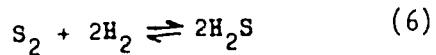
The formation of magnesium sulfate, therefore, is no primary reaction. The influence of a gaseous reducing agent upon the

Card 2/4

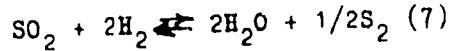
SOW/76-4-2-5/40

Investigation of the Thermal Decomposition of Magnesium Sulfite

thermal decomposition of magnesium sulfite does not show a considerable difference from the reaction in nitrogen atmosphere. Upon reduction of gaseous substances sulfuretted hydrogen appears:



The formation of elementary sulfur increases in hydrogen atmosphere on the effect of catalysts, as is shown in the equation:



Thus, in the presence of bauxite in the hydrogen current, a large amount of the sulfur of magnesium sulfite turns into elementary sulfur and sulfuretted hydrogen and a small amount only into sulfur dioxide during the decomposition of magnesium sulfite in the temperature range of 400-500°. Therefore no magnesium thiosulfate is formed. The experimental results are of theoretical and practical importance. The method can be used for the production of elementary sulfur. There are 1 figure, 4 tables, and 5 Soviet references.

Card 3/4

AMIROVA, S.A.; PECHKOVSKIY, V.V.; KURMAYEV, R.Kh.

Solubility of vanadium trioxide in fused sodium and potassium chlorides. Zhur. neorg. khim. 9 no.5:1229-1231 My '64.  
(MIRA 17:9)

1. Permskiy politekhnicheskiy institut.

PECHKOVSKIY, V.V.; MAL'TSEVA, T.G.

Interaction of calcium oxide with sulfur dioxide under  
reducing conditions. Zhur. prikl. khim. 37 no.2:240-  
246 F '64. (MIHA 17:9)

1. Permskiy politekhnicheskiy institut.

L 27267-65 EWT(m)/EPF(c)/EWA(d)/T/EWP(t)/EWP(b) pr-4 IJP(c) JD/JG/WB

ACCESSION NR: AP4011442

S/0076/64/038/001/0108/0114

29  
25

B

AUTHORS: Amircva, S.A. (Perm'); Pechkovskiy, V.V. (Perm'); Prokhorova, V.G. (Perm'); Zhebeleva, T.V. (Perm'); Lezhneva, A.A. (Perm')

TITLE: Oxidation of manganese-vanadium spinel by oxygen

SOURCE: Zhurnal fiz. khim. v. 38, no. 1, 1964, 108-114

TOPIC TAGS: manganese vanadium spinel, manganese vanadium spinel oxidation, spinel decomposition, manganese metavanadate, manganese pyro vanadate

ABSTRACT: The oxidative annealing of manganese-vanadium spinel was investigated from 0 to 1000°C, using thermographic analysis simultaneously with x-ray, crystallooptic and chemical methods. The first stage of the oxidation is the chemisorption of oxygen on the surface of the spinel grains and the formation of a solid solution. Decomposition of the spinel at both high and low temperatures proceeds according to the following equations:  $2\text{MnO}\cdot\text{V}_2\text{O}_3 + 2\text{O}_2 = \text{Mn}_2\text{V}_2\text{O}_7 + \text{V}_2\text{O}_5$ ;  $\text{Mn}_2\text{V}_2\text{O}_7 + \text{V}_2\text{O}_5 = 2\text{Mn}(\text{VO}_3)_2$ . The

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L 27267-65

ACCESSION NR: AP4011442

products are acid soluble and almost insoluble in water. The melting points of manganese metavanadate and pyrovanadate are given as 805 and 1023°C respectively. Small additions of potassium chloride intensify the rate of oxidation. Orig. art. has: 2 equations, 10 figures and 1 table.

ASSOCIATION: Pernskiy politekhnicheskiy institut (Perm Polytechnical Institute)

SUBMITTED: 05Mar63

ENCL: 00

SUB CODE: MM

NR REF SOV: 005

OTHER: 004

Card 2/2

26039  
S/137/61/000/007/006/072  
A060/A101

18.3200

AUTHORS: Pechkovskiy, V. V., Zvezdin, A. G.

TITLE: Study of sulfating cobalt concentrate ash in a fluidized bed

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 7, 1961, 19, abstract 70136  
("Uch. zap. Permsk. un-t", 1959, v. 13, no. 3, 119-123)

TEXT: The rate and completeness of sulfating of an ash Co compound with SO<sub>2</sub> - air mixtures as a function of temperature, SO<sub>2</sub> concentration in the gas, granule size, initial degree of sulfation of the Co and sulfide S concentration in the ash, duration of roasting in the fluidized bed reactor, are studied. Factory ash, obtained by the roasting of Co concentrate from the Pyshminskoye Mine Administration, was used for the investigation. The initial degree of sulfation of Co in the ash constituted 64 and 9.0 percent. The mean total content of Co in the ash was 0.705 percent. The ash was granulated by means of sulfide alkalies. As result of the investigation it is recommended to submit the granulated ash to an additional sulfation by mixtures of SO<sub>2</sub> and O<sub>2</sub> in a fluidized bed furnace. The additional sulfation of Co in the ash proceeds at a remarkable rate at 550°C and attains a maximum at 700°C after which it decreases. An X

Card 1/2

26039  
S/137/61/000/007/006/072  
A060/A101 X

Study of sulfating cobalt concentrate ...

increase in  $\text{SO}_2$  concentration in the gaseous phase exerts a positive effect on the sulfation process. The sulfation of ash in the fluidized bed proceeds at high intensity during the first 15 - 20 min, after which the rate of the process slows down. The optimal size of ash granules is 1.0 - 2.0 mm. As the sulfide content of the ash increases to S > 8 percent the rate of sulfate formation falls sharply and the maximum Co sulfation is displaced to a lower temperature zone.

N. Pleteneva

[Abstracter's note: Complete translation]

Card 2/2

SOV/80-32-5-5/52

(2)

AUTHOR:

Pechkovskiy, V.V.

TITLE:

The Study of the Factors Affecting the Process of Sulfide Formation of Metal Oxides by Sulfur Dioxide

PERIODICAL:

Zhurnal prikladnoy khimii, 1959, Vol 32, Nr 5, pp 966-969 (USSR)

ABSTRACT:

The sulfide formation of metal oxides by sulfur dioxide in a reducing atmosphere is studied here. The method of the experiment and the characteristics of the used substances are given in Ref. 5. The sulfurous anhydride was diluted by dry, pure nitrogen. As carbon-containing materials blood coal, semi-coke, coals from various layers birch charcoal, etc. were used. The particle size of the metal oxides and the carbon-containing materials was less than 0.078 mm. If the concentration of the sulfurous anhydride decreases from 100 to 20%, the degree of sulfidation is gradually reduced. At a decrease from 20 to 5% the sulfidation is reduced sharply. At less than 5% the sulfide formation is slow and incomplete. At 850°C and a gas concentration of 20% the used carbon-containing materials may be arranged according to their effect in the following decreasing series: charcoal, blood coal, semi-coke, Kuznetsk coal, coke. The sulfide formation is decreased by a

Card 1/2

SOV/80-32-5-5/52

The Study of the Factor Affecting the Process of Sulfide Formation of Metal Oxides  
by Sulfur Dioxide

dense crystal structure and a surface of low reactivity. An addition of manganese oxide  $Mn_2O_3$  increases the sulfide formation, ferrous bauxite reduces it. If cobalt and zinc oxides are burnt with Kuznetsk coal, sulfide formation decreases, if the coal particles are larger than 0.25 mm. With charcoal the decrease starts with sizes larger than 1 mm. The rate of sulfidation is highest in the first 20-30 min, then slows down sharply.

There are: 6 tables and 5 Soviet references.

ASSOCIATION: Permskiy gosuniversitet imeni A.M. Gor'kogo (Perm" State University  
imeni A.M. Gor'kogo)

SUBMITTED: March 5, 1958

Card 2/2

5.4220

75651  
SOV/80-32-10-10/51AUTHOR: Pechkovskiy, V. V.

TITLE: Reaction of Barium and Calcium Carbonates With Sulfur Dioxide in the Presence of a Reducing Agent

PERIODICAL: Zhurnal prikladnoy khimii, 1959, Vol 32, Nr 10, pp 2189-2193 (USSR)

ABSTRACT: This is a study of the effect that temperature, percentage and type of carbon-containing material, and  $\text{SO}_2$  concentration in the gas have in the above reaction on the yield of sulfites, sulfates, and sulfides on a  $\text{CaCO}_3$  and  $\text{BaCO}_3$  basis. Evidence of the following reactions was found:

$$\text{MeCO}_3 \rightleftharpoons \text{MeO} + \text{CO}_2 \quad (1)$$
$$\text{MeO} + \text{SO}_2 \rightleftharpoons \text{MeSO}_3 \quad (2)$$
$$4\text{MeSO}_3 + 2\text{SO}_2 = 4\text{MeSO}_4 + \text{S}_2 \quad (3)$$
$$\text{SO}_2 + \text{C} \rightleftharpoons \frac{1}{2}\text{S}_2 + \text{CO}_2 \quad (4)$$
$$\text{CO}_2 + \text{C} \rightleftharpoons 2\text{CO} \quad (5)$$
$$2\text{MeO} + \text{S}_2 + \text{C} = 2\text{MeS} + \text{CO}_2 \quad (6)$$
$$\text{MeS} + 2\text{SO}_2 = \text{MeSO}_4 + \text{S}_2 \quad (7)$$

Card 1/4

Reaction of Barium and Calcium Carbonates With  
Sulfur Dioxide in the Presence of a Reducing  
Agent

75651  
SOV/80-32-10-10/51

I. Decomposition of  $\text{CaCO}_3$ . Starting at  $700^\circ$ , with increasing temperature, the yield of  $\text{CaS}$  increases, and the yield of  $\text{CaSO}_3$  and  $\text{CaSO}_4$  decreases; this is because the rate of reactions (1), (4), and (6) increases with temperature, while (2) goes to the left and (4) to the right. Above  $900^\circ$ , provided there is enough C,  $\text{CaS}$  is the main sulfur-containing product. Substitution of  $\text{CaO}$  for  $\text{CaCO}_3$  has little effect on the reaction under study. Within the  $650$  to  $750^\circ$  range, the yield of  $\text{CaS}$ ,  $\text{CaSO}_4$ , and  $\text{CaSO}_3$  is greater, all other conditions being equal, than in the case of  $\text{CaCO}_3$ , because reaction (1) limits the rate of formation of  $\text{CaS}$ ,  $\text{CaSO}_4$  and  $\text{CaSO}_3$ ; at  $\geq 750^\circ$ , (1) is no longer limiting. II. Decomposition of  $\text{BaCO}_3$ . Similarly, the  $\text{BaS}$  yield increases with temperature, while the curves of temperature vs percentage

Card 2/4

Reaction of Barium and Calcium Carbonates With  
Sulfur Dioxide in the Presence of a Reducing  
Agent

75661  
SOV/80-32-10-10/51

conversion to  $\text{BaSO}_4$  and to  $\text{BaSO}_3$  exhibit maxima. Above  $1,000^\circ\text{C}$ , provided enough C is present,  $\text{BaS}$  will be the main product. The  $\text{BaS}$ , and especially the  $\text{BaSO}_4$  and  $\text{BaSO}_3$  yields are greater, all other conditions being equal, than the  $\text{CaS}$ ,  $\text{CaSO}_4$ , and  $\text{CaSO}_3$  yields; also, the percentage conversion peaks of  $\text{BaSO}_4$  and  $\text{BaSO}_3$  are shifted in the direction of higher temperatures in comparison with the corresponding Ca compounds. This is due to the greater thermal stability of the Ba compounds. Effect of %C: All other conditions being equal, for both Ba and Ca carbonates, the sulfide yield decreases and the sulfate yield increases with decreasing %C because the rates of (3) and (7) increase, while the rates of (4) and (6) decrease. Effect of type of carbon-containing material: At  $900^\circ\text{C}$ , coke reactivity is slight, but at  $> 1,000^\circ\text{C}$ , it approaches the reactivity of charcoal. Effect of  $\text{SO}_2$  concentration: Owing to an

Card 3/4

Reaction of Barium and Calcium Carbonates With Sulfur Dioxide in the Presence of a Reducing Agent 75661 SOV/80-32-10-10/51

increase in the rate of (2) to (4), (6), and (7), an increase in  $\text{SO}_2$  concentration will raise the yield of Ca and Ba sulfur-containing compounds; the net percentage of sulfur-containing Ba products increases, largely owing to an increase in  $\text{BaSO}_4$  concentration, while the BaS percentage actually decreases. There are 4 tables; and 9 Soviet references.

ASSOCIATION: Perm' State University (Permskiy gosudarstvennyy universitet)

SUBMITTED: December 10, 1958

Card 4/4

PECHKOVSKIY, V.V.; AMIROVA, S.A.; POLTONYANSHECHIKOVA, M.I.

Investigation of preliminary slag roasting on the vanadium recovery process. Izv. vys. ucheb. zav.; tsvet. met. 3 no.3:97-101 '60. (MIRA 14:3)

1. Permskiy gosudarstvennyy universitet, Kafedra tekhnologii neorganicheskikh veshchestv.  
(Vanadium-Metallurgy) (Slag)

AMIROVA, S.A.; PECHKOVSKIY, V.V.; PROKHOROVA, V.G.; POLOTNYANSHCHIKOVA, M.I.

Study of the oxidation roasting of converter slags for the extraction  
of vanadium. Izv.vys.ucheb.zav.; khim.i khim.tekh. 3 no.6:1056-1061  
'60. (MIRA 14:4)

1. Permskiy politekhnicheskiy institut, kafedra tekhnologii  
neorganicheskikh veshchestv.  
(Vanadium) (Slag)

PECHKOVSKIY, V.V.; AMIROVA, S.A.; KETOV, A.N.

Reduction of iron, zinc, manganese, magnesium and calcium sulfates  
by hydrogen and carbon. Uch. zap. Perm. gos. un. 17 no.1:3-  
14 '60. (MIRA 14:11)

(Reduction, Chemical)  
(Sulfates)

PECHKOVSKIY, V.V., KETOV, A.N.

Thermography in studying topochemical reactions. Uch. zap. Perm.  
gos. un. 17 no.1:15-34 '60. (MIRA 14:11)  
(Thermal analysis)  
(Chemical reactions)

PECHKOVSKIY, V.V.; AMIROVA, S.A.; KETOV, A.N.

Intensification of the reduction of sulfates by adding sodium  
and potassium carbonates. Uch. zap. Perm. gos. un. 17 no.1:45-  
54 '60. (MIRA 14:11)

(Sulfates)  
(Reduction, Chemical)

LOZHIN, A.F.; PECHKOVSKIY, V.V.; SUBOCHEVA, N.L.

Effect of additives of some potassium compounds on the reduction  
process of barites. Uch. zap. Perm. gos. un. 17 no.1:55-60 '60.  
(MIRA 14:11)

(Barite)  
(Potassium compounds)

AMIROVA, S.A.; PECHKOVSKIY, V.V.; KAMEKO, B.S.; STEPANOVA, A.F.

Investigating methods for using pickling solutions. Uch. zap.  
Perm. gos. un. 17 no.1:61-72 '60. (MIRA 14:11)  
(Metals—Pickling)

"APPROVED FOR RELEASE: 06/15/2000

CIA-RDP86-00513R001239810018-8

PECHKOVSKIY, V.V.; AMIROVA, S.A.; KAMEKO, G.F.; POLOTNYANSHCHIKOVA, M.I.

Investigating the granulation and firing of vanadium slag with  
additives. Uch. zap. Perm. gos. un. 17 no.1:83-90 '60.  
(MIRA 14:11)

(Vanadium)

APPROVED FOR RELEASE: 06/15/2000

CIA-RDP86-00513R001239810018-8"

PECHKOVSKIY, V.V.

Sulfating cobalt and zinc compounds with sulfur dioxide. Uch.  
zap. Perm., gos. un. 17 no.1:91-95 '60. (MIRA 14:11)  
(Sulfur dioxide)  
(Zinc)  
(Cobalt)

PECHKOVSKIY, V.V.

Study of the reduction roasting of zinc and magnesium sulfates.  
Zhur.prikl.khim. 33 no.5:1025-1030 My '60. (MIRA 13:7)

1. Permskiy gosudarstvennyy universitet imeni A.M. Gor'kogo.  
(Zinc sulfate) (Magnesium sulfate) (Reduction, Chemical)

PECHKOVSKIY, V.V.; KETOV, A.N.

Reduction of barium sulfate with carbon monoxide at high temperatures.  
Zhur. prikl. khim. 33 no.8:1719-1723 Ag '60. (MIRA 13:9)  
(Barium sulfate) (Carbon monoxide)

"APPROVED FOR RELEASE: 06/15/2000

CIA-RDP86-00513R001239810018-8

PECHKOVSKIY, V.V.; KETOV, A.N.

Thermal decomposition of zinc sulfite. Zhur. prikl. khim. 33  
no.8:1724-1729 Ag '60. (MIRA 13:9)  
(Zinc sulfite)

APPROVED FOR RELEASE: 06/15/2000

CIA-RDP86-00513R001239810018-8"

PECHIKOVSKII, V. V.; AMIROVA, S.A.; KAMEKO, B. S.

Study of the ways of utilising hydrolysis acid. Zhur. prikl. khim.  
33 no.9:1976-1981 S '60. (MIRA 13:10)

1. Permskiy gosudarstvennyy universitet imeni A.M. Gor'kogo  
(Sulfuric acid)

*S. T. M.*  
PECHKOVSKIY, V. V., Doc Tech Sci, "INVESTIGATION OF  
CALCINATION OF SULFUR-CONTAINING COMPOUNDS." LENINGRAD,  
1961. (MIN OF HIGHER AND SEC SPEC ED RSFSR, LENINGRAD  
ORDER OF LABOR RED BANNER TECHNOL INST IN LENSOVET).  
(KL, 3-61, 212).

165

S/081/62/000/011/036/057  
E071/E192

AUTHORS: Ketov, A.N., and Pechkovskiy, V.V.  
TITLE: A study of the reactivity of carbonaceous materials using differential thermal analysis  
PERIODICAL: Referativnyy zhurnal, Khimiya, no.11, 1962, 502, abstract 11 M 34. (Sb. nauchn. tr. Permsk. politekhn. in-t, no.10, 1961, 65-70).  
TEXT: Using a Kurnakov pyrometer, heating rate curves of carbonaceous materials, heated in a stream of air and CO<sub>2</sub>, filtering through the substance studied, were obtained. The heating rate was 25 °C/min., the fineness of materials 0.25-0.5 mm, the rate of gas flow 5 litres/hour. For charcoal the exothermic reaction with air was observed at 292 °C and for graphite at 348 °C. The temperatures of the beginning of interaction with CO<sub>2</sub> were found: charcoal 721 °C, semicoke 856 °C, coke 854 °C. A dilution of CO<sub>2</sub> with nitrogen does not change the temperature of the endothermic peak, but decreases its value. Saturation of charcoal with solutions of Na<sub>2</sub>CO<sub>3</sub> and K<sub>2</sub>CO<sub>3</sub>

Card 1/2

S/137/62/000/005/031/150  
A006/A101

AUTHORS: Amirova, S. A., Pechkovskiy, V. V., Prokhorova, V. G., Polotnyanshchikova, M. I.

TITLE: Roasting of granulated and moistened vanadium-containing charges in an enlarged laboratory furnace

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 5, 1962, 18, abstract 50108 ("Sb. nauchn. tr. Permsk. politekhn. in-t", 1961, no. 10, 111 - 119)

TEXT: The authors studied optimum conditions of roasting granulated V-charges and the possibility of combining granulation, drying and roasting of moistened V-charges in an enlarged rotating tubular furnace. For this purpose, mixtures consisting of converter slag, sylvinitite, and refuse slime, were granulated and roasted. Best results were obtained when roasting granules of 2 - 5 mm fraction at 850°C with addition of sylvinitite ( $n = 0.5$ ) and 5% refuse. The degree of V extraction was 94 - 95%. Roasting of moistened, freshly prepared granules at 850°C makes it possible to extract up to 95% V. Roasting of the charge with simultaneous granulation of the material in the furnace is possible, the charge

Card 1/2

Roasting of granulated and...

S/137/62/000/005/031/150  
A006/A101

is granulated to 94 - 95%,  $V_2O_5$  extraction is 95% at  $950^{\circ}C$  and 4 - 4.5 hours roasting time. Best results in simultaneous granulation, drying and roasting are obtained with a charge consisting of slag with addition of sylvinit (n = 0.5) and 5% refuse, with 10.5% moisture at a slope angle of the furnace of  $1^{\circ}30'$ . When employing such methods of roasting V-containing slags, V extraction increases up to 94 - 95%. There are 7 references.

G. Svodtseva

[Abstracter's note: Complete translation]

Card 2/2

S/137/62/000/005/032/150  
A006/A101

AUTHORS: Amirova, S. A., Pechkovskiy, V. V., Prokhorova, V. G., Polotnyan-shchikova, M. I., Derendyayeva, M. P.

TITLE: Preliminary oxidizing as a means of raising the degree of vanadium extraction from converter slags

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 5, 1962, 18, abstract 5G109 ("Sb. nauchn. tr. Permsk. politekhn. in-t", 1961, no. 10, 121 - 129)

TEXT: Studies were conducted of the optimum conditions for roasting V-charges composed of previously oxidized slag and alkaline admixtures. All experiments were made on an enlarged laboratory rotary furnace 2,500 mm long with 100 mm inner diameter. Initial material was converter slag of the following composition (in %): V<sub>2</sub>O<sub>5</sub> 13.5; MnO 3.8; MgO 0.95; Fe<sub>disp</sub> 3.1; FeO 37.9; TiO<sub>2</sub> 8.2; SiO<sub>2</sub> 31.4; Cr<sub>2</sub>O<sub>3</sub> 9.1; CaO 1.1; Al<sub>2</sub>O<sub>3</sub> 2.04. KCl and commercial sylvinitie containing NaCl 74.5% and KCl 22%, were employed as alkaline admixtures. The molar ratio, in composing the charge, of the alkaline admixture to

Card 1/2

S/137/62/000/005/032/150  
A006/A101

Preliminary oxidizing as a means of...

$V_2O_5$  of the slag was 1 or 0.5. Oxidized slag was obtained by roasting the initial slag in an enlarged laboratory furnace for 6 hours at 850 - 880°C; it contained 11.45%  $V_2O_5$ . During its leaching out in the laboratory with  $H_2SO_4$  of 7% concentration, 91.5% V were extracted into the solution. The prepared and thoroughly mixed charges were placed into the furnace. The duration of roasting was regulated by changing the slope angle and the rotation speed of the furnace pipe. It was found that roasting of a charge of previously oxidized slag and sylvinitic permits up to 94 - 95% V extraction at 800°C during 7 hours; 91% V is extracted in the form of water soluble compounds. The addition to the charge of waste slag from the Chusovo Metallurgical Plant in a 10% amount, reduces caking and increases the degree of V extraction. There are 5 references.

G. Svodtseva

[Abstracter's note: Complete translation]

Card 2/2

8/137/62/000/005/030/150  
A006/A101

AUTHORS: Amirova, S. A., Pechkovskiy, V. V., Prokhorova, V. G., Derendyayeva,  
M. P.

TITLE: Developing a new technology of extracting vanadium from converter  
slags

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 5, 1962, 17 - 18, abstract  
5G107 ("Sb. nauchn. tr. Perms. politekhn. in-t", 1961, no. 10,  
131 - 137)

TEXT: On the model of a generalized laboratory unit optimum conditions  
were studied for the following stages of a technological system for extracting  
V from slag: oxidizing roasting of crushed converter slag without admixtures;  
hot acid lixiviation of the roasted slag, and precipitation of  $V_2O_5$  from the  
solutions obtained. The experiments were made with converter slags of the fol-  
lowing chemical composition (in %):  $V_2O_5$  13.5; FeO 37.9;  $Cr_2O_3$  9.1; MnO 3.8;  
 $TiO_2$  3.2; CaO 1.1; MgO 0.95;  $SiO_2$  34.4;  $Al_2O_3$  2.0 and  $Fe_{met}$  3.1. It was  
found that optimum conditions for roasting non-granulated slag are as follows:

Card 1/2

LOZHIN, A.F.; PECHKOVSKIY, V.V.; SUBOCHEVA, N.L.

Formation of acid-soluble barium compounds in the reduction  
roasting of barite. Izv. vys. ucheb. zav.; khim. i khim.  
tekhn. 4 no. 2:242-246 '61. (MIRA 14:5)

1. Permskiy gosudarstvennyy universitet im. A.M. Gor'kogo.  
Kafedra tekhnologii neorganicheskikh veshchestv.  
(Barium salts) (Barite)

LOZHIN, A.F.; PECHKOVSKIY, V.V.; SUBOCHEVA, N.L.

Reduction firing of granulated barite concentrates. Izv.vys.  
ucheb.zav; khim.i khim.tekh. 4 no.5:832-836 '61. (MIRA 14:11)

1. Permskiy gosudarstvennyy universitet, kafedra tekhnologii  
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carbon. Zhur.prikl.khim. 38 no.6:1199-1206 Je '65.

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CIA-RDP86-00513R001239810018-8

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Thermochemical transformations of chromium and manganese chlorides.  
Zhur. neorg. khim. 9 no.9:2059-2065 S '64.

(MIRA 17:11)

L 1066-65 ENG(j)/DWT(m)/EPF(c)/EPR/EWP(l) Pr-4/Ps-4 SSD/AS(mp)-3/  
AEDC(a)/RAEM(i)/AFETR/AFWL/ASD(m)-5/ASD(a)-5/ESD(t) JD/JW/jG

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AUTHOR: Pechkovskiy, V. V.; Zvezdin, A. G.

TITLE: Kinetics of vanadium trioxide oxidation

SOURCE: Kinetika i kataliz, v. 5, no. 3, 1964, 424-429

TOPIC TAGS: vanadium trioxide, oxidation, kinetics, vanadium pentoxide, process mechanism, autocatalysis, diffusion, activation energy, additive, oxidation acceleration

ABSTRACT: The oxidation of  $V_2O_3$  was investigated to determine whether the process is autocatalytic and whether it is possible to establish the mechanism of the process with formal kinetic equations. The experiments were run in apparatus described by O. Kadlets, M. N. Dubinin (Izv. AN SSSR, Otd. khim. n. 1031, 1956) wherein the samples were heated to the desired temperature in a closed system, then the timer was engaged as air was admitted (circulating at different rates with the help of a pump). The oxidation product is  $V_2O_5$ . At all temperatures investigated (260-460) the oxidation starts rapidly, decreases sharply at first and then more slowly. It was established that the maximum on the rate of reaction

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ACCESSION NR: AP4041063

duration of heating curve, which was sometimes considered an indication of the existence of autocatalysis, is in the case of  $V_2O_3$  merely an indication of the increase in reaction rate occurring while heating the materials to the test temperature. Arrhenius formal kinetics equations describe the kinetics of the process in the 300-480°C range. When the degree of oxidation is over 60% and the temperature is above 460°C the effects of diffusion inhibition (diffusion of oxygen through the product layer) become strongly apparent. The calculated apparent energy of activation of the oxidation of  $V_2O_3$  with atmospheric oxygen varies from 29-37 kcal. Addition of KCl has no accelerating effect on the oxidation. Orig. art. has: 2 tables, 7 figures and 3 equations.

ASSOCIATION: Pernskiy politekhnicheskiy institut (Pernsk Polytechnical Institute)

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Card 2/2

AMIROVA, S.A.; PECHKOVSKIY, V.V.; PROKHOROVA, V.G.; OSTROVSKAYA, T.V.;  
LEZHNEVA, A.A. (Perm')

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1. Permskiy politekhnicheskiy institut.

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zav.; khim. i khim. tekhn. 6 no.6:991-996 '63. (MIRA 17:4)

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AMIROVA, S.A.; PECHKOVSKIY, V.V.; VARSKOY, E.N.; TYULENEVA, G.Ye.

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Drying and oxidative roasting of granular vanadium slag in a laboratory fluid-bed furnace. Izv.vys.ucheb.zav.;khim.i khim.tekh. 6 no.4:625-630 '63. (MIRA 17:2)

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PECHKOVSKIY, V.V.; AMIROVA, S.A.; VOROB'YEV, N.I.

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(Slag)

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(Mammal training) (Tools)